Fiberoptic Probe For Hemodynamic Spectroscopy in Freely Moving Mice Shows Cerebrovascular Dysregulation In Alzheimer's and Glioma Models

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Abstract:

Significance: Cerebral vascular reactivity is critical parameters of brain homeostasis in health and disease, but the investigational value of brain oxymetry is diminished by anesthesia and mechanical fixation of the mouse skull.

Aim: We needed to reduce the physical restrictivity of hemodynamic spectroscopy to enable cancer and Alzheimer's disease (AD) studies in freely-moving mice.

Approach: We combined spectroscopy, spectral analysis software and a magnetic, implantable device to measure vascular reactivity in unanesthetized, freely-moving mice. We measured cerebral blood volume fraction (CBVF) and oxygen saturation (S_{O_2}).

Results: CBVF and S_{O_2} demonstrated delayed cerebrovascular recovery from hypoxia in an orthotopic xenograft glioma model and we found increased CBVF during hypercapnia in a mouse model of AD compared to wild-type littermates.

Conclusions: Our optomechanical approach to reproducibly getting light into and out of the brain enabled us to successfully measure CBVF and S_{O_2} during hypercapnia in unanesthetized freely-moving mice. We present hardware and software enabling oximetric analysis of metabolic activity, which

provides a safe and reliable method for rapid assessment of vascular reactivity in murine disease models as well as CBVF and S_{O_2} .

Keywords: fiberoptic spectroscopy, cerebral oxymetry, Alzheimer's disease mouse model, glioma, cerebrovascular reactivity

1. Introduction

1.1 Importance of cerebrovascular reactivity

Cerebral perfusion is tightly regulated in response to ischemia (reactive hyperaemia) or the metabolic demand associated with increased local neuronal activation (functional hyperaemia). Vascular reactivity is therefore a fundamental property of blood vessels and enables rapid adjustment of vascular flow to meet the metabolic needs of brain tissue, and is coordinated by the interaction of neurons, glia, endothelium and pericytes(1). Alteration of this response impairs the ability to provide sufficient oxygen and nutrients to the brain and leads to disease states(2). Impaired vascular reactivity has been implicated in Alzheimer's disease (AD)(2, 3). AD patients show decreased cerebral hemoglobin oxygen saturation (S_{O_2}) during verbal fluency tasks(4) and a failure of cerebral blood flow (CBF) increase during visual stimulus tests(5). Mouse models of AD also exhibit cerebrovascular dysregulation during whisker stimulation or vasodilator treatment(6). In addition to AD, brain tumors -- particular gliomas -- show high intratumoral vascular heterogeneity with regions of vascular hyperplasia and avascular necrosis interspersed(7). In gliomas, microvascular proliferation in response to tumor hypoxia is closely associated with malignant progression and poor survival(3).

1.2 Limitations of Current Technology

Several diagnostic modalities, including functional Magnetic Resonance Imaging (fMRI)(8), functional Near-Infrared Spectroscopy (fNIRS)(9) and visible light fiberoptic spectroscopy(10), have been used to measure vascular reactivity in animal studies and patients. In animal studies, these techniques require anesthesia in order to minimize movement artifacts, which prevents analysis of cerebral perfusion during behavioral tasks. In addition, anesthesia is known to significantly uncouple neural and vascular responses(11), leading to modified cerebrovascular function. The ability to study changes in cerebral S_{O_2}

and cerebral blood volume fraction (CBVF) without anesthesia could greatly contribute to a more accurate understanding of oxygen needs and cerebrovascular changes in various neurological disorders.

1.3 Novel fiberoptic method for measuring S_{O_2} and CBVF in freely moving animals

In order to overcome these limitations, we developed a device and method for measuring S_{O_2} and CBVF in freely moving animals using fiberoptic spectroscopy. S_{O_2} depends on the balance of blood flow and oxygen consumption by the tissue. S_{O_2} and CBVF, which are both unit-less, together characterize the oxygen content in the tissue. Compared to fMRI, fiberoptic spectroscopy is inexpensive and portable, and offers higher temporal resolution. Compared to fNIRS, visible light spectroscopy penetrates only superficially considering the optical scattering of brain tissue (Fig. 1), offering measurement in a banana-shaped volume between a source fiber which illuminates the brain surface and a radially-displaced detector fiber (Fig. 2). With deeper implantation of the probe fibers, cerebrovascular measurements of deep brain regions, such as the hypothalamus or amygdala, in response to complex behavioral tasks will be possible with our approach. We developed a connection between the mouse brain and the spectrometer consisting of an implanted brain probe (~1 cm fiber) and a spectrometer connector (~2 m fiber). The device (Fig. 3a) fiber-optically coupled the brain to the light source and to the spectrometer when magnetically coupled but could be magnetically detached so that the probe could be surgically implanted prior to surgical recovery and subsequent measurement during normal behavior (Fig. 3b,c). After recovery and attachment, spectral measurements were analyzed by least-square fitting to light transport theory (Fig. 3d) to specify the instantaneous S_{O_2} and CBVF. These values varied slightly (Fig. 3e) during normal behavior and $CBVF > 0$ indicated proper attachment reliably.

2. Materials and Methods

2.1 System and computational method of fiberoptic measurement

A white light source (HL-2000-HP, Ocean Optics, Dunedin FL) illuminated the cerebral cortex through an illumination fiber bundle of 8 fibers (FG105UCA, Thorlabs, Newton, NJ) in 7-around-1 mated to a solid core fiber in the tip (FT400UMT, Thorlabs, Newton, NJ), while a identical detection fiber tip and bundle directed the measured diffuse reflectance to a spectrometer (HDX, Ocean Optics, Dunedin FL). A less sensitive spectrometer (e.g. HR-2000) was used for some experiments but the integration time is much slower (~0.3s) so we prefer the high temporal resolution (~10Hz) of the HDX instead. Our system was controlled through Matlab (Mathworks, Natick, Massachusetts) software by a laptop computer running Windows 10. Spectra were fit for real-time readout of $CBVF$ and S_{O_2} on the perioperative laptop during experimentation. The graphic user interface is provided in the supplementary materials.

2.2 Spectral Calibration

Spectra were calibrated using measurements of a 99% reflectance standard (AS-011XX-X60, Labsphere, North Sutton, NH) while holding the fiber probe 3 cm from the standard. The dark noise of the spectrometer was also measured (dark). Calibrated spectral measurements $M(\lambda)$ were calculated as a function of wavelength (λ) from the raw tissue measurements $R(\lambda)$ and the calibration measurements $C(\lambda)$ according to **Equation 1**.

$$M(\lambda) = \frac{R(\lambda)}{C(\lambda)} \quad M(\lambda) = \frac{R(\lambda) - \text{dark}}{C(\lambda) - \text{dark}} \quad (1)$$

where K is a scaling factor, essentially the ratio of collection efficiency of embedded fibers to collection efficiency of fiber probe held above standard. The measured optical spectrum was dependent on the spectra of the light source and the detector sensitivity but these factors were cancelled by the normalization (Equation 1). A predicted spectrum (M_p) was created by multiplying a predicted R_p based on diffusion theory by a scaling factor (K), $M_p = KR_p$. Equations 2-12 specify the calculation of R_p , provided here to facilitate reproducibility of M_p and implement a method we previously developed (10,

12), which uses the diffusion theory expression of Farrell, Patterson, and Wilson (13-16). The use of this analytical approach (Equations 2-12) was valid, since the 2.5 mm source-detector separation was large compared to the transport mean free path [$1/(\mu_a + \mu_s') = 1.26$ mm at 630 nm], and the scattering dominated over absorption.

$$r_i = 0.6681 + 0.0636 \times n + 0.7099 / n - 1.4399 / n^2 \quad (2)$$

$$A = \frac{(1 + r_i)}{(1 - r_i)} \quad (3)$$

$$z_0 = \frac{1}{\mu_a(\lambda) + \mu_s'(\lambda)} \quad (4)$$

$$D = \frac{z_0}{3} \quad (5)$$

$$\Delta = \sqrt{\frac{D}{\mu_a(\lambda)}} \quad (6)$$

$$r_1 = \sqrt{z_0^2 + r^2} \quad (7)$$

$$r_2 = \sqrt{(z_0 + 4AD)^2 + r^2} \quad (8)$$

$$\mu_{eff} = \frac{1}{\Delta} \quad (9)$$

$$c = z_0 \left(\mu_{eff} + \frac{1}{r_1} \right) \times \frac{e^{-\frac{r_1}{\Delta}}}{r_1^2} \quad (10)$$

$$d = (z_0 + 4AD) \times \left(\mu_{eff} + \frac{1}{r_2} \right) \times \frac{e^{-\frac{r_2}{\Delta}}}{r_2^2} \quad (11)$$

$$M_p = \frac{c + d}{4\pi} \quad (12)$$

The fiber separation (distance between the source fiber tip and the detector fiber tip) was $r \approx 0.25$ cm, and the refractive index mismatch between glass ($n_{\text{glass}} = 1.52$) and tissue ($n_{\text{tissue}} = 1.35$) was $n = n_{\text{glass}}/n_{\text{tissue}} = 1.13$. $\mu_a(\lambda)$ and $\mu_s'(\lambda)$ were the absorption and reduced scattering coefficients of the tissue, respectively. The spectral absorption coefficient, $\mu_a(\lambda)$, was calculated as a weighted sum of the absorption spectra of oxygenated whole blood ($\mu_{a_oxy}(\lambda)$), deoxygenated whole blood ($\mu_{a_deoxy}(\lambda)$), and water ($\mu_{a_water}(\lambda)$), according to Equation 13:

$$\mu_a(\lambda) = CBVF \times [S_{O_2} \times \mu_{a_oxy}(\lambda) + (1 - S_{O_2}) \times \mu_{a_deoxy}(\lambda)] + W \times \mu_{a_water}(\lambda) \quad (13)$$

2.3 Calculation of Fitting Parameters to Quantify S_{O_2} and CBVF

Spectra were analyzed in the range 540 to 650 nm by least-squares fitting. The difference between M (Eq. 1) and $M_p = KR_p$ (see Eq. 12) was minimized by adjusting M_p using a multidimensional unconstrained nonlinear minimization method (Nelder-Mead), the `fminsearch()` in Matlab. Fitting adjusted 3 variables: $CBVF$, S_{O_2} and K where $CBVF = 1$ specifies 150 g/L hemoglobin. W , the fractional tissue water content, was fixed at an assumed value of 0.78, which has been reported for mouse cortex (17). Absorption due to water was minimal. For example, at 650 nm, low $CBVF = 0.01$ and $S_{O_2} = 0.5$, the absorption coefficient in the tissue due to the presence of blood is $\mu_{a,\text{blood}} = CBVF \times S_{O_2} \times \mu_{a,\text{oxy}} + CBVF \times (1 - S_{O_2}) \times \mu_{a,\text{deoxy}} = 0.1103 \text{ cm}^{-1}$. $\mu_{a,\text{water}} = 0.0025 \text{ cm}^{-1}$, which is 44-fold lower than $\mu_{a,\text{blood}}$.

The absorption of the tissue due to blood was specified by the fitting described above while the scattering optical properties (scattering coefficient $\mu_s(\lambda)$ [cm^{-1}] and scattering anisotropy $g(\lambda)$ [-]), which were constant across all experiments, were spectral functions (λ) fit to the previous measurements of fresh human brain tissue (18). Thus $\mu_s(\lambda)$ and $g(\lambda)$ were used as interpolated functions of wavelength (**Equations 14-19**) that, in combination yielded the reduced scattering coefficient, $\mu_s'(\lambda) = \mu_s(\lambda) \times (1 - g(\lambda))$ used in **Equation 4**. These optical properties agree reasonably with the literature (18, 19), such as our previous data (18) that can be fit for the reduced scattering coefficients of gray and white matter as shown in Fig. 1C.

2.4 Brain optical scattering properties

Fig. 1 shows the brain scattering properties specified by the spectral data at 5 wavelengths from Yaroslavsky et al. 2002 (18). The Fig. 1a and Fig. 1b show the spectral behavior of the scattering coefficient (μ_s) and the anisotropy of scattering (g). The combination of μ_s and g yields the reduced scattering coefficient ($\mu_s' = \mu_s(1 - g)$) shown in Fig. 1c for white and gray matter. Scattering properties of whole tissue were set as a weighted-sum based on a 1:2 ratio of white to grey matter, as reported (20) for mouse cortex.

$$\begin{aligned} \text{Tissue_}g &= \text{WhiteMatter_}g * \frac{1}{3} + \text{GrayMatter_}g * \frac{2}{3} \\ \text{Tissue_}\mu_s &= \text{WhiteMatter_}\mu_s * \frac{1}{3} + \text{GrayMatter_}\mu_s * \frac{2}{3} \end{aligned}$$

(14)

(15)

The $\mu_s'(\lambda)$ spectrum was assumed when using least-square fitting of the data to specify K, CBVF and S_{O_2} . After spectral fitting, the values for CBVF and S_{O_2} were output as a function of time (e.g. **Fig. 1e**) over time periods where spectroscopic measurements were made on mice under various experimental conditions. These optical properties were also used to model the photon transport in the mouse brain.

2.5 *Fiberoptic probe with surgically implantable, magnetically-coupling tip*

The connection between the mouse brain and the spectrometer consisted of an implanted brain probe (~1 cm fiber) and a spectrometer connector (~2 m fiber). The connector (Fig. 3a) fiber-optically coupled to the illumination source and the spectrometer and opto-magnetically coupled to the brain probe, which fiber-optically coupled the brain.

Plastic connectors were 3D-printed using a 3D Systems ProJet printer and Visijet M3 Crystal material. Mating surfaces were 3D-printed facing upward (opposite the print bed) to minimize warp and maximize surface flatness. The plastic connector design was optimized to align and hold (press fit) two ceramic ferrule cannulas (CF440-10, Thor Labs, Newton NJ) and two cylinder magnets (D12-N52, K&J Magnetics, Plumsteadville PA). During assembly, the connector's flat surface was pressed against glass, and the ferrules and magnets were inserted and also pressed against the glass for alignment (friction fit). The male side connector was fabricated using a similar process of press fitting the cannulae and magnets, but the assembly was pressed against an alignment tool (see Supplemental Materials), instead of the flat glass plate. The connector was then mated to its probe with magnetic attachment, and pre-polished fibers (FT400UMT, Thor Labs, Newton NJ) were inserted into the ferrules and affixed using UV-cure adhesive (#68, Norland Products, Cranbury NJ). After dry, the protruding fibers were clipped

and polished to a length of 0.5 mm protruding from the cannulas using a custom polishing tool set (see Supplemental Fig. 1) that held the probe such that the fibers were normally incident on standard fiber polishing film.

In order to maintain alignment, so that the fibers in the brain probe and spectrometer connector mated precisely, the brain probe surface was printed with 6 small 4-sided pyramids while the connector surface was printed with complementary mating holes. This design configuration enabled secure mating of the two connecting parts during measurements.

Finished fiber probes were visually inspected for polish level and analyzed through a series of signal and noise tests using the spectrometer and the light source targeted on human skin. Human skin was chosen as a biological tissue calibration target because it was readily accessible to the researchers. A threshold for optical performance in satisfactorily-manufactured fiber probes was set as a 60-1 signal-to-noise ratio. The signal was the spectrally summed diffuse remittance with the fiber probe touching skin. The noise, which consisted of light “leaking” directly from the emission fiber of the connector and entering the detection fiber of the connector (ie. not passing through the brain probe), was the sum of the spectrally summed signal measured while pointing the probe into the air in a dark room.

2.6 Analysis of cerebrovascular reactivity of freely moving mice

30 mice, over two years during technical development of our probe, were retrofitted with implantable probe tips and measured during normal behavior. 7 month-old C57BL/6 mice (Jackson Laboratory, Bar Harbor, ME) were used to measure basal cerebrovascular dynamic properties of the cortex while freely moving. The AD model mice studied were 13 month-old Tg6799 (Jackson Laboratory) double transgenic mice overexpressing both human amyloid precursor protein (APP) gene with KM670/671/NL, V717I, and I716V mutations and human presenilin 1 (PSEN1) harboring M146L and L286V mutations under the Thy1 promoter (21). Wildtype (WT) littermates were used as controls. Only

male mice were used in experiments. All mice were maintained in The Rockefeller University's Comparative Biosciences Center and treated in accordance with protocols approved by Rockefeller University's Institutional Animal Care and Use Committee (IACUC). After anesthesia with intraperitoneal injection of ketamine/xylazine (100 mg/kg and 10 mg/kg, respectively), mice were mounted on a stereotactic frame. After the removal of hair over the scalp, the scalp was cleaned with sterile alcohol and Betadine three times. We performed a minimal sized midline scalp incision (~5 mm). We then drilled two small holes in the skull using a 0.7 mm diameter stainless steel micro drill burr (Fine Science Tools). The coordinates of the two holes were Bregma= -1 mm, lateral= 1.5 mm and Bregma=-3.5 mm, lateral=1.5 mm. Brain probes were fitted to the holes and immobilized using dental cement. Brain probes were fitted to each hole and immobilized using dental cement. The skin incision was sutured and animals were monitored throughout the recovery period. A week after surgery, mice were placed into a chamber (22.8 cm x 20.3 cm x 15 cm) and we measured cerebral perfusion in the freely moving state. For the hypercapnia study, we injected 10%CO₂/10% O₂/80%N₂ mixture in to the induction chamber and measured changes in *CBVF* and *S_{O2}* during and after hypercapnia.

2.7 Analysis of cerebrovascular reactivity in mouse model of glioma

A separate group of mice were used for the glioblastoma model at the New York University. These mice were housed within NYU School of Medicine's Animal Facilities. Brain tumor implantations were performed according to protocols approved by NYU's IACUC, as previously described (22). Briefly, 6-8 week-old NOD.SCID (NOD.CB17-Prkdcscid/J, 001303, Jackson Laboratory, Bar Harbor, ME) male mice were anesthetized with intraperitoneal injection of ketamine/xylazine (100 mg/kg and 10 mg/kg, respectively). They were then mounted on a stereotactic frame (Harvard Apparatus). A midline skin incision was made. A high-speed drill was used to drill a small hole in the calvaria 2 mm off the midline and 2 mm anterior to coronal suture. Five µl of a suspension of U87 cells (100,000 cells/µl) were injected through a Hamilton syringe (1 µl/min, Harvard apparatus, needle pump) into the left frontal lobe through the drilled hole. The injection needle was left in place for an additional 5 min after the injection.

was completed to prevent backflow. The skin incision was sutured and animals were monitored throughout the recovery period. U87-MG cells for production of intracranial xenografts were cultured in Dulbecco's Minimal Essential Media (DMEM, Life Technologies) supplemented with 10% FBS and non-essential amino acids. Previous studies with U87-MG cells in our laboratory showed large tumor formation 2 weeks after injection (22). Two weeks after the injection of cells, terminally ill animals were prepared for fiberoptic measurement. After anesthesia and fitting into the stereotactic frame as described above, we drilled an additional hole in the contralateral frontal calvaria, using a high-speed drill. The diameter of the 2 holes matched the caliber of the probe. After cleaning the area with ethanol wipes, the probes were fitted to each window and immobilized using dental cement. Animals were removed from the stereotactic frame for fiberoptic measurements. We first measured CBVF and S_{O_2} at the resting state for 30 sec. To analyze the cerebrovascular response during hypoxia, we blocked the nostrils for 30 sec with a wet cotton swap, and measured CBVF and S_{O_2} during hypoxia. After nostril blockage release, we continued CBVF and S_{O_2} recordings during reperfusion. This procedure was repeated once, then animals were euthanized following the approved protocol. The skull was opened and the presence of the glioma in situ under the probe was confirmed by a neuropathologist.

3. Results

3.1 Analysis of cerebrovascular reactivity in freely moving C57BL/6 and AD mice

Fiber spectroscopic measurement of CBVF and S_{O_2} in mixed arterial and venous blood the cortex of freely moving 7-month old C57BL/6 mice yielded values of ~1.4% and ~55% respectively (Fig. 4a). To investigate the change in CBVF and S_{O_2} of the cortex during hypercapnia in the mouse cortex, we placed mice in an insulated chamber and increased CO_2 concentration up to 10%. When hypercapnia was induced after 70 seconds of baseline measurement by increasing CO_2 concentration, we found an increase in CBVF, which is a typical hyperaemic response to increased CO_2 through vasodilation. S_{O_2} decreased slowly but normalized rapidly after stopping CO_2 injection and releasing of CO_2 at 280 sec

(Fig. 4b), while CBVF increased rapidly before returning to a level that was less than the baseline measurement.

To investigate the cerebrovascular properties of an AD (Tg6799) (21) mouse, we measured the baseline of CBVF and S_{O_2} of the cortex of freely moving 13 month-old Tg6799 mice and wild-type (WT) littermates and also analyzed the difference of vascular reactivity between Tg6799 mice and WT littermates during and after hypercapnia using fiber optic spectroscopy (Fig. 4c,d). While there is no difference in baseline of CBVF and S_{O_2} between the two groups, the increase of CBV of the cortex of AD mice during hypercapnia was significantly higher than that of WT littermates (Fig. 4e). In addition, although not statistically significant, the time to baseline of CBVF of AD mice was also delayed in comparison to WT littermates (Fig. 4f).

3.1 Analysis of cerebrovascular reactivity in glioma mouse model

To demonstrate broad utility we analyzed cerebrovascular response to hypoxia in an orthotopic xenograft mouse model generated with human U87 glioblastoma (GBM) cells. Since patients with brain tumors show impaired cerebral vasoreactivity (23-25) and hyperproliferation of immature blood vessels lacking smooth muscle cells limiting perfusion in malignant glioma (3), an orthotopic xenograft glioma mouse model is a good platform to test our new system. Two weeks after surgical implantation of the fiberoptic probe on the skull of control mice (sham injected) and mice orthotopically injected with human U87 GBM cells, we induced hypoxia for 30 seconds and measured the S_{O_2} and CBVF change directly from brain parenchyma. In sham mice, during hypoxia for 30 sec, cerebral S_{O_2} decreased but normalized within 30 seconds after hypoxic release (Fig. 4g). We observed the same pattern of hypoxia and rapid normalization at each cycle of obstruction/release as expected for a normal cerebrovasculature. Comparing the AD recovery time to that of the glioma model mice to the example mouse recovery time for the glioma model, the glioma mouse had a recovery of more than 30 seconds to recover their normal S_{O_2} (Fig. 4h). This result suggests that the cerebrovasculature of glioma has impaired vasoreactivity and reduced ability to restore perfusion after hypoxia.

4. Discussion

Fiberoptic spectroscopy can assess the S_{O_2} and the $CBVF$ in the brains of freely moving mice when implemented with a magnetically coupling tip. The present study showed that our fiberoptic method detected changes in freely moving mice and measured cerebrovascular impairment in a mouse model of GBM. Fig. 4 shows typical S_{O_2} and $CBVF$ values in GBM and background mouse strains. Though our measurements of freely-moving and glioblastoma mice were separate, and we measured only normal mice moving freely and disease mice anesthetized, we hope to measure freely behaving mice of disease-mouse models. S_{O_2} in cerebral blood of control mice was 45-60%, which is comparable to previous studies in the cortex of anesthetized rodents (26, 27). $CBVF$ of 6-8 week old NOD.SCID mice was ~2.0% (Fig. 4a), and $CBVF$ of 7 month old C57BL/6 mice was 1.4% (Fig. 4b). This range from 1% to 2% in our results is slightly less than the range reported in the literature: 2.1% reported by Synchrotron Radiation Quantitative Computed Tomography in the parietal cortex (28), 2.03-3.14% reported by MR in the cortex (29-31), 2.0-2.4% reported by multiphoton laser scanning microscopy and in capillary-rich cerebral cortex (32). We measured $CBVF$ of 7-month old mice, instead of young rodents (4-8 weeks old), which other studies mainly used (29-32).

This fiberoptic method may improve understanding the mechanisms of impaired vascular reactivity in neurologic disorders and brain tumors. Our vasculature and hemodynamic measurements raise the question as to whether the observed phenomena are consequences of the pathology or more fundamental changes that are required for disease progression. We hope to explore these temporal measurements in future work, potentially combining $CBVF$ and S_{O_2} with time-differential mathematics to access perfusion and oxygen consumption parameters. Understanding the extent of impaired vasoreactivity in brain pathologies may someday enable novel diagnostic and prognostics without need for ex-vivo studies.

Supplementary Materials: The 3D models to print our device, the commercial component part list and the operating system software to replicate our work are described in the supplemental documentation and provided at the following location for download:

Fig. 1. Spectral scattering coefficient (A) and scattering anisotropy (B) from Yaroslavsky et al. 2002 (18). The fits are combined to yield the spectrum of the reduced scattering coefficient (C), where a is the μ_s at 650 nm and b is the scattering power.

Fig. 2. Monte Carlo simulation outputs a map of the of the surgically implanted probe

The light distribution is shown within the murine skull and brain tissue at 650 nm wavelength that is both delivered by the source fiber and collected by the collection fiber. The distribution is the joint

probability of delivering to and collecting from each voxel. $f_{\text{delivered}} \times f_{\text{collected}}$, which has units of $[\text{cm}^{-2}] \times [\text{cm}^{-2}]$ or $[\text{cm}^{-4}]$, where f denotes fluence $[\text{cm}^{-2}]$. The figure plots $\log_{10}(f_{\text{delivered}} \times f_{\text{collected}})$.

Fig. 3. Instrumentation and typical readout for Fiberoptic cerebral oximetry in freely moving mice.

(A), 3D model of magnetic-coupled probe. 6 male/female mating pyramid pairs secure the detachable, implantable brain probe into a linking fiberoptic with mating female pyramid components to enable fiberoptic alignment. (B), surgically-implanted brain probe. (C), brain probe connected to spectrometer. (D), spectral fitting in <0.25 seconds yielding instantaneous measurements of the percentages of CBVF and SO_2 . (E), Typical time-course measurement over normal behavior. The graph shows cerebral blood volume fraction (blue) times 100 such that a y-axis value of 1 means 1% blood volume fraction and the oxygen saturation fraction (red), which is the fraction of hemoglobin molecules bound to oxygen within the cerebral probed volume.

Fig. 4: In vivo spectroscopy results

a, Basal cerebrovascular properties of the cortex of a freely moving, 7 month-old mouse. **b**

Cerebrovascular responses in 7 month-old freely moving mice were obtained using fiberoptic spectroscopy during hypercapnia. Data are representative of three independent experiments. **c**

& **d**, Cortical vascular reactivity of Tg6799 13 month-old mice (**c**) and WT littermate (**d**) during hypercapnia were analyzed using fiberoptic spectroscopy. Hypercapnia

was induced at 70 sec by increasing CO₂ concentration up to 10 % and was kept until 300 sec. Data are representative of four independent

experiments. **e**, The increase of cerebral blood volume fraction (CBVF) during hypercapnia was significantly higher in Tg6799 mice compared to WT littermate (p<0.05, n = 4). **f**, The time to baseline of CBV after hypercapnia was also delayed in Tg6799 mice compared to WT littermate, but it is not statistically significant (p = 0.06, n = 4). **g**, Hypoxic conditions were held from 30 – 60 sec and from 80 – 110 sec in 2 month-old mouse. The graph shows cerebral blood volume fraction (blue) and the oxygen saturation fraction (red). **h**, Delayed Cerebrovascular recovery from hypoxia in a NOD.SCID 2 month-

old mouse with intracranial U87 GBM xenograft was delayed compared to a sham mouse. Bar graphs represent mean $\pm$ s.e.m. of ≥ 3 independent experiments, and was statistically analyzed by performing a two-tailed unpaired t-test.

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